

Phenolic compounds of Argan tree, Argania spinosa (endemic species of South Western Morocco)

Saïda Tahrouch, Abdelhakim Hatimi, Sylvie Rapior, Laurence Mondolot,

Annick Gargadennec, Claude Andary

► To cite this version:

Saïda Tahrouch, Abdelhakim Hatimi, Sylvie Rapior, Laurence Mondolot, Annick Gargadennec, et al.. Phenolic compounds of Argan tree, Argania spinosa (endemic species of South Western Morocco). TOJSAT: The Online Journal of Science and Technology, 2011, 1 (4), pp.17-23. hal-02201606

HAL Id: hal-02201606 https://hal.umontpellier.fr/hal-02201606v1

Submitted on 31 Jul 2019

HAL is a multi-disciplinary open access archive for the deposit and dissemination of scientific research documents, whether they are published or not. The documents may come from teaching and research institutions in France or abroad, or from public or private research centers. L'archive ouverte pluridisciplinaire **HAL**, est destinée au dépôt et à la diffusion de documents scientifiques de niveau recherche, publiés ou non, émanant des établissements d'enseignement et de recherche français ou étrangers, des laboratoires publics ou privés.

PHENOLIC COMPOUNDS OF ARGAN TREE, ARGANIA SPINOSA (ENDEMIC SPECIES OF SOUTH WESTERN MOROCCO)

Saïda Tahrouch¹, Abdelhakim Hatimi¹, Sylvie Rapior², laurence Mondolot², Annick Gargadennec² and Claude Andary²

¹Laboratoire de Biotechnologies Végétales, Département de Biologie, Faculté des Sciences, Université Ibnou Zohr, B.P. 8106-Dahkla Agadir, Maroc.

² Laboratoire de Botanique, Phytochimie et Mycologie, UMI-CNRS (UPR 9056, CEFE), Faculté de Pharmacie, Université Montpellier I, F-34060 Montpellier cedex 2.

e-mail: tahrouch@hotmail.com

Abstract: Argania spinosa (L.) Skeels (Sapotaceae) is an endemic tree located mainly in south-western of Morocco. The argan tree plays medicinal, ecological and socioeconomic roles in this area. The fruit of A. spinosa has oil-producing kernels with a high unsaturated fatty acid content. The argan oil is greatly used in food and cosmetic products. Kernel, pulp of fruit and trunk have also been studied for sterols, triterpenes and saponins. Our goal in this study is to investigate the leaves for phenolic compounds by HPLC in 90 specimens of argan tree from three localities in Souss Massa area (south-western Morocco). Quantification and histolocalisation of phenolic components, i. e. flavonoids and condensed tannins (molecules well known for their broad spectrum of biological activities) in the three localities were carried out using chromatographic and spectroscopic methods combined to histochemical technics. Flavonol glycosides were quantified by HPLC from argan leaves. The main flavonol glycoside was myricitrin. The content of myricetin derivatives was higher than the quercetin derivative content. With regard to chemotaxonomy, four flavonol glycosides seem to be good markers for this species as they were detected by HPLC in 90 specimens of argan tree from the 3 localities. The histochemical studies of the different parts of A. spinosa (leaves, stems and thorns) have shown a high concentration of myricetin derivatives in the peripheral tissues, this cell localisation of the flavonoids could explain the Argan tree adaptation to aridity.

Key words: *Argania spinosa* - Flavonol glycosides – Condensed tannins – Histochemistry.

INTRODUCTION

Argania spinosa (L.) Skeels (*Sapotaceae*) is an endemic and medicinal tree (Bellakhdar J., 1997) located mainly in south-western Morocco. The argan tree plays ecological and socioeconomic roles in this area (Boukhobza and Pichon-Prum, 1988).

The fruit of *A. spinosa* has oil-producing kernels with a high unsaturated fatty acid content (Maurin *et al.*, 1992). The argan oil is greatly used in food (Huyghebaert and Hendrickx, 1974) and cosmetic products (Pierre Fabre Patent). Whereas kernel, pulp of fruit and trunk have been studied extensively for sterols, triterpenes and saponins (Farines *et al.*, 1984; Charrouf *et al.*, 1991, 1992; Maurin, 1992; Nerd *et al.*, 1994; Oulad-Ali *et al.*, 1996), relatively little is known about chemistry and histochemistry of the leaves, the stems and the thorns (Tahrouch *et al.*, 1998; Tahrouch *et al.*, 2000).

In order to understand both vigour and resistance of this endemic plant to an arid habitat, the leaves were investigated for phenolic compounds by HPLC in 90 specimens of argan tree from three localities. Quantification of phenolic components, i. e. flavonoids and condensed tannins in three localities were carried out using chromatographic and spectroscopic methods combined to histochemical techniques.

MATERIAL AND METHODS

The leaves of *Argania spinosa* were collected in the Ademine reserve (**A**) 60 m, Ait Baha (**AB**) 500 m and Immouzzer (**I**) 900 m, Agadir (Morocco). Voucher specimens were deposited in the Herbarium of Laboratoire des Symbiotes Racinaires et de Biochimie Végétale in Agadir. Dry leaves were extracted three times with MeOH/H₂O (4/1) at room temperature.

Histochemistry

Sections of leaves (45-60 μ m thickness) were cut with a cryostat microtome (Frigocut 2800 E) operating at -20°C and examined using either a light microscope or an epi-fluorescence microscope (Nikon Optiphot) with two filter sets: UV filter set with 365 nm excitation and a 400 nm barrier filter. Flavonoid compounds were detected using Neu's reagent (Neu, 1956). Sections were immersed into the reagent for 1 min and then observed by epi-fluorescence (Dai *et al.*, 1995). DMCA (4-dimethylaminocinnamaldehyde) reagent was used to locate condensed tannins (Feucht *et al.*, 1986). Stained sections were observed with a light microscope.

Quantification of flavonoids and condensed tannins

HPLC was carried out using an isocratic mobile phase (Acetonitrile/MeOH/H₂O, 2/8/15) running through Nucleosil C18 (250 x 4 mm, 5 μ m particle size). UV-Visible data were recorded using a Photodiode Array Detector coupled to HPLC system. Flavonoids of argan leaves were quantified using myricitrin as internal standard at 350 nm.

Quantitative determination of condensed tannins were carried out by UV and visible spectrophotometry according to Mc Murrough and Mc Dowell (1978).

RESULTS

A. spinosa was investigated for condensed tannins and flavonoids, molecules well known for their broad spectrum of biological activities, (Di Carlo *et al.*, 1999). Flavonol glycosides were quantified by HPLC from argan leaves (figure 1). The main flavonol glycoside (Tahrouch *et al.*, 2000) was myricitrin [2]. The content of myricetin derivatives [2, 4] was higher (\cong 20 mg.g⁻¹ D. W.) than the quercetin derivative [1, 3] content (\cong 8 mg.g⁻¹ D. W.).

Compounds	Quercitrin	Myricitrin	Hyperoside	Myricetin-3-O-galactoside [4]
	[1]	[2]	[3]	
Rt ^a	15.2	8.8	10.1	6.8
Q ^b	5.3±0.4	16.8±1.4	2.5±0.2	3.3±0.3

^a retention time (minutes); ^b quantity (mg.g⁻¹D. W. \pm s.e.)

With regard to chemotaxonomy, these four molecules seem to be good markers for this species as they were detected by HPLC in 90 specimens of argan tree from the 3 localities : A, AB and I (tables 1, 2 and 3).

These results showed that there is a close relationship between A and AB. The amount of flavonoids in these 2 localities (A and AB) is higher than in I. A and AB are located in arid areas.

It might be advisable to combine analytical and histological methods. With histochemistry we are able to localise *in situ* the flavonoids by Neu's reagent, which give a bright orange-yellow fluorescence

under UV. Neu's reagent is a borate salt that forms complex with certain groups of phenolic compounds giving them specific fluorescence (doc. 1). Condensed tannins detected histochemically by using DMCA reagent give a blue coloration under white light (doc. 2).

The histochemical studies of the different parts of *A. spinosa* (leaves, stems and thorns) have shown a high concentration of myricetin derivatives in the peripheral tissues particularly in epidermis while condensed tannins were mainly deposited in the cortex and palisade mesophyl.

The high content of total flavonoids in specimens of *A. spinosa* that located in localities A and AB, could play a protective role in the expression of tolerance to UV-radiations as showed by Lois (1994) in *Arabidopsis thaliana (Brassicaceae)*. Olsson *et al.* (1998) explained that flavonoids afforded a protective role not only through the absorption of UV-radiations, especially in the epidermal layers, but also through a selective increase after UV-B irradiation that happens for flavonoids which possess an additional hydroxyl group in the B-ring of the flavonoid skeleton such as quercetin and myricetin derivatives that we identified from *A. spinosa* leaves.

The increase of phenolic compound biosynthesis in plant under UV-radiation and during periods of water stress and nutrient deficiency (Keller and Hrazdina, 1998) might be an adaptation phenomena (Gershenzon, 1984). Indeed, the biosynthesis of flavonoids in plants is enhanced in response to changes in the external environment (Cooper-Driver and Bhattacharya, 1998). According to this hypothesis, plants that normally occupy arid area and infertile habitats, such as argan trees, could have continuously high levels of phenolic constituents. The high amount of phenolic components and their localisation in the peripheral tissues might contribute to understand the relationship between accumulation of polyphenols and adaptation of argan tree to his area.

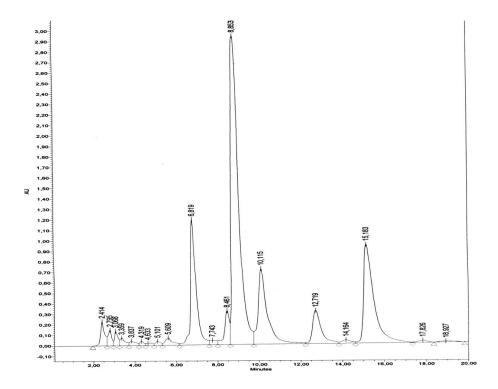


Figure 1 : HPLC of methanolic extract of Argan leaves.

components		M2	M3	M4	M5	M6	M7	M8	M9	M10
	M1									
retention time	2,41	2,80	5,60	6,81	8,46	8,85	10,11	12,71	14,16	15,18

M4	:	Myricetin 3-O-galactoside (D)
M6	:	Myricitrin (B)
M7	:	Hypéroside (C)
M10	:	Quercitrin (A)
M5	:	Myricetin derivatives
M8	:	Quercetin derivatives
The o	the	r flavonoids (M1, M2, M3, and M9) were not identified.

Table 1 : Quantification of flavonoids (M1-M10) and condensed tannins (M11) in Admine reserve ($mg.g^{-1}$ D.W.).

	M1	M2	M3	M4 (D)	M5	M6 (B)	M7 (C)	M8	M9	M10 (A)	M11
1	1,460	1,426	0	2,740	0	10.037	2,318	1,517	1,134	2,214	59,90
2	1.623	1.800	1.055	2.081	1,143	15,484	1,054	1.118	1.729	5,892	44.00
3	1,327	1,545	1,093	1,875	0	8.670	2,368	1,438	1,503	4,674	139,35
4	1,536	1,330	0	0	0	7,190	0	0	0	1,698	47,35
5	1,504	1,368	1,089	2,646	1,062	9,635	1,945	1,404	1,182	2,454	78,75
6	1,248	1,219	1,169	3,488	0	15,129	2,330	1,187	1,566	4,734	69,55
7	1,288	1,281	0	1,608	0	10,392	0	1,042	1,244	3,162	81,30
8	1,334	1,341	1,096	4,330	0	12,842	3,184	1,804	1,252	3,115	72,90
9	1,228	1,266	0	1,643	0	6,312	0	1,040	1,066	1,955	60,35
10	1,447	1,724	0	2,231	0	12,562	2,499	1,305	1,354	3,924	60,35
11	1,338	1,547	1,085	2,607	1,147	18,753	1,039	1,205	1,596	5,219	90,55
12	1,356	1,526	1,163	2,763	1,340	14,220	3,522	1,539	1,608	5,494	68,30
13	1,486	1,592	1,224	3,098	1,305	19,165	2,678	1,375	1,901	7,860	144,30
14	1,256	1,345	1,142	2,443	1,063	9,150	2,742	1,649	1,467	4,131	32,70
15	1,768	1,833	0	4,107	0	28,365	4,159	1,429	2,819	14,381	91,55
16	1,285	1,349	0	2,891	1,097	11,639	2,739	1,559	1,664	5,669	52,40
17	1,594	1,666	1,468	5,402	0	21,789	3,439	1,632	1,318	3,686	103,50
18	1,267	1,378	1,038	2,858	0	14,132	2,421	1,451	1,323	3,318	65,80
19	1,547	1,662	1,414	4,776	1,327	26,907	1,067	1,587	1,522	4,724	63,70
20	1,546	1,513	1,135	4,204	1,202	21,866	2,999	1,300	1,734	5,550	69,60
21	1,115	1,196	0	1,579	0	8,061	0	1,080	1,156	2,398	45,40
22	1,580	1,698	1,090	2,947	1,079	16,483	3,196	1,464	1,956	7,040	78,10
23	1,534	1,564	1,196	4,648	1,246	27,859	3,920	1,695	2,894	11,972	87,85
24	1,158	1,269	1,050	2,627	0	11,538	2,051	1,218	1,309	3,347	111,10
25	1,196	1,383	1,317	2,096	0	12,266	2,410	1,201	1,646	5,044	102,00
26	1,311	1,396	1,270	3,391	0	20,390	0	1,235	1,574	4,705	39,90
27	1,617	1,570	1,465	2,337	1,208	15,267	2,446	1,438	1,851	6,541	61,15
28	1,325	1,450	1,354	2,286	1,040	10,558	2,804	1,636	1,446	4,291	61,55
29	1,498	1,466	1,434	3,618	1,113	18,156	2,744	1,502	1,522	4,914	72,60
30	1,557	1,534	1,302	2,676	1,091	17,152	2,577	1,430	1,608	5,267	70,05

Table 2 : Quantification of flavonoids (M1-M10) and condensed tannins (M11) in Aït Baha	i reserve (mg.g ⁻¹
D.W.).	

	M1	M2	M3	M4 (D)	M5	M6 (B)	M7 (C)	M8	M9	M10 (A)	M11
1	1,459	1,4264	0	2,740	0	10,036	2,317	1,516	1,133	2,2136	60,70
2	1,622	1,7996	1,054	2,080	1,143	15,483	1,054	1,117	1,729	5,8924	62,00
3	1,326	1,5452	1,092	1,874	0	8,669	2,367	1,437	1,502	4,674	50,95
4	1,536	1,33	0	0	0	7,190	0	0	0	1,6976	58,15
5	1,504	1,3684	1,088	2,646	1,062	9,635	1,945	1,404	1,182	2,4536	99,00
6	1,248	1,2192	1,169	3,487	0	15,129	2,330	1,187	1,565	4,7344	70,90
7	1,287	1,2812	0	1,608	0	10,392	0	1,042	1,244	3,162	50,10
8	1,334	1,3412	1,095	4,330	0	12,842	3,184	1,804	1,251	3,1148	53,50
9	1,227	1,2656	0	1,642	0	6,311	0	1,039	1,064	1,9548	51,80
10	1,447	1,7244	0	2,230	0	12,562	2,499	1,304	1,353	3,924	112,2
11	1,338	1,5472	1,085	2,607	1,146	18,753	1,038	1,205	1,595	5,2192	49,65
12	1,356	1,5256	1,163	2,763	1,340	14,219	3,522	1,539	1,608	5,494	72,15
13	1,485	1,592	1,223	3,098	1,305	19,165	2,678	1,374	1,901	7,8596	72,15
14	1,256	1,3452	1,141	2,442	1,063	9,1504	2,741	1,649	1,466	4,1308	60,30
15	1,768	1,8328	0	4,107	0	28,364	4,158	1,429	2,819	14,3808	130,30
16	1,284	1,3492	0	2,890	1,096	11,639	2,738	1,559	1,663	5,6692	43,30
17	1,594	1,666	1,467	5,401	0	21,789	3,438	1,632	1,318	3,6856	81,50
18	1,267	1,378	1,038	2,858	0	14,132	2,421	1,450	1,322	3,318	47,10
19	1,546	1,6624	1,414	4,775	1,327	26,906	1,067	1,586	1,522	4,724	59,45
20	1,546	1,5128	1,134	4,204	1,202	21,865	2,999	1,300	1,733	5,55	64,10
21	1,115	1,196	0	1,578	0	8,061	0	1,080	1,156	2,3976	31,40
22	1,580	1,6984	1,090	2,947	1,078	16,482	3,195	1,464	1,955	7,04	70,05
23	1,534	1,5644	1,196	4,647	1,246	27,858	3,920	1,695	2,894	11,9716	44,55
24	1,158	1,2688	1,049	2,626	0	11,538	2,050	1,218	1,309	3,3468	30,55
25	1,196	1,3828	1,316	2,096	0	12,266	2,409	1,201	1,646	5,0444	37,80
26	1,310	1,3964	1,269	3,391	0	20,390	0	1,234	1,574	4,7048	58,15
27	1,616	1,57	1,464	2,337	1,208	15,267	2,446	1,438	1,850	6,5408	64,95
28	1,325	1,45	1,354	2,285	1,040	10,558	2,803	1,636	1,446	4,2912	57,30
29	1,498	1,4656	1,434	3,617	1,112	18,156	2,744	1,501	1,522	4,9144	61,95
30	1,557	1,534	1,302	2,676	1,090	17,152	2,576	1,429	1,607	5,2672	103,00

	M1	M2	M3	M4 (D)	M5	M6 (B)	M7 (C)	M8	M9	M10 (A)	M11
1	0	0	0	1,270	0	4,234	1,834	1,205	1,248	2,682	9,80
2	0	0	0	0	0	5,142	1,340	0	1,153	2,208	17,95
3	0	0	0	1,326	0	6,211	1,960	1,334	1,227	2,862	20,40
4	0	0	0	0	0	1,865	0	1,247	0	0	8,63
5	0	0	0	1,600	0	3,102	1,812	1,145	0	1,588	14,70
6	0	0	1,225	2,238	0	6,707	2,258	1,401	1,186	2,478	26,30
7	0	0	0	2,107	0	7,148	3,265	1,446	1,531	4,302	26,50
8	1,073	0	0	2,019	0	8,776	2,946	1,662	1,493	4,170	28,55
9	0	0	1,200	1,078	0	2,554	1,399	1,120	1,114	2,080	19,60
10	1,064	1,130	0	2,096	0	8,979	2,496	1,505	1,333	3,231	33,85
11	1,130	1,150	1,836	3,552	0	15,080	5,596	2,871	2,285	7,092	55,25
12	0	1,096	0	1,716	0	7,125	1,966	1,273	1,259	2,859	29,60
13	0	0	0	3,013	0	9,233	2,688	1,714	1,311	2,642	47,75
14	0	1,138	0	1,930	0	6,905	2,670	1,352	1,442	3,643	26,50
15	1,458	1,470	1,434	3,086	0	17,116	4,060	2,266	2,401	6,754	64,10
16	1,097	1,042	1,836	2,861	0	6,348	2,474	1,867	1,289	2,756	30,20
17	1,150	1,096	0	1,609	0	6,210	1,888	1,284	1,502	4,257	39,80
18	0	1,050	0	2,746	0	10,995	3,082	1,556	1,994	7,053	31,60
19	1,440	1,377	0	1,466	0	7,841	1,881	1,291	1,442	3,740	97,55
20	1,260	1,362	0	3,098	0	11,747	4,166	1,962	1,875	5,566	49,45
21											12,65
22	1,232	1,476	0	5,232	0	13,406	5,368	2,608	1,992	5,956	55,25
23	1,207	1,255	0	1,328	0	3,524	1,517	1,150	1,086	1,962	54,45
24	1,059	0	1,045	3,211	0	10,314	3,852	1,904	1,740	5,054	36,85
25	1,170	1,246	0	2,663	0	7,971	3,128	1,739	1,529	4,084	51,10
26	1,247	1,344	1,495	2,847	0	10,469	3,654	1,967	1,856	5,230	45,25
27	0	0	0	1,900	0	6,533	2,755	1,489	1,520	3,834	19,60
28	1,503	1,462	0	2,370	0	17,876	3,230	1,742	2,158	6,254	59,10
29	1,041	1,141	0	0	0	3,987	0	0	1,302	2,928	38,10
30	0	1,127	0	2,153	0	6,617	2,544	1,458	1,242	2,586	27,75

Table 3 : Quantification of flavonoids (M1-M10) and condensed tannins (M11) in Immouzer reserve $(mg.g^{-1} D.W.)$.

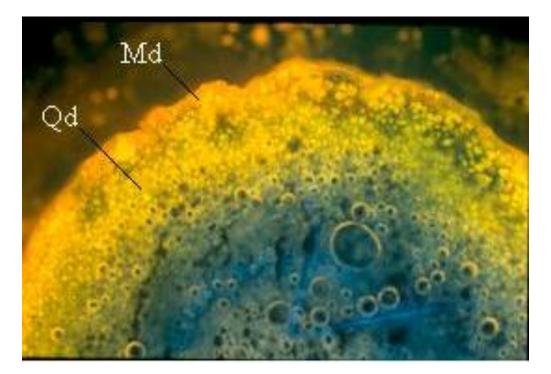


Figure 2 : The histochemical study from the stem of *A. spinosa* showed a high content of myricetin derivatives (Md) in the peripheral tissues. Qd : Quercetin derivatives.

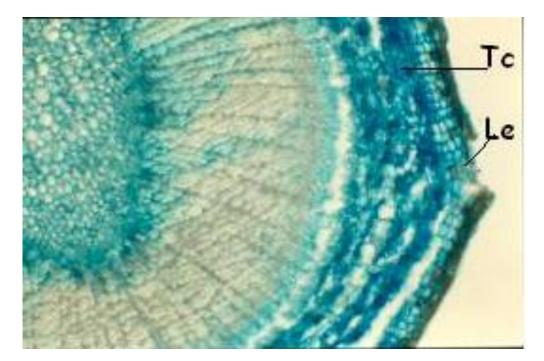


Figure 3 : The histochemical studie from the stem of *A. spinosa* showed a high concentration of condensed tannins (Tc) mainly deposited in the cortex. Le : lenticel.

REFERENCES

Bellakhdar, J. (1997).- La pharmacopée marocaine traditionnelle (Médecine arabe ancienne et savoirs populaires). Ed. Ibis Press, Saint-Etienne, 776p.

Boukhobza, M., Pichon-Prum, N. (1988). L'arganier, ressource économique et médicinale pour le Maroc. *Phytotherapy*, **27**, 21-26.

Charrouf, Z., Wieruszeski, J.M., Fkih-Tetouani, S., Leroy, Y., Charrouf, M., Fournet, B. (1992). Triterpenoid saponins from *Argania spinosa*. *Phytochemistry*, **31**, 2079-2086.

Charrouf, Z., Fkih-Tetouani, S., Charrouf, M., Fournet, B. (1991). Triterpenes and sterols extracted from the pulp of *Argania spinosa* (L.) *Sapotaceae*. *Plantes Médicinales et Phytothérapie*, **25**, 112-117.

Cooper-Driver, G.A., Buattacharya, M. (1998). Role of phenolics in plant evolution. *Phytochemistry*, **49**, 1165-1174.

Dai, G. H., Andary, C., Mondolot-Cosson, L., Boubals, D. (1995). Histochemical responses of leaves of *in vitro* plantlets of *Vitis spp*. to infection with *Plasmopara viticola*, *Phytopathology*, **85**, 149-154.

Di Carlo, G., Mascolo, N., Izzo, A.A., Capasso, F., (1999). Flavonoids: Old and new aspects of a class of natural therapeutic drugs. *Life Sciences*, **65**, 337-353.

Fabre P. Cosmétique S. A. Lipidic extract of argan fruit and its use in cosmetology. P. Hatinguais, M. T. Trebosc & R. Belle, (Patent) FR 83-16740.

Farines M., J. Soulier, M. Charrouf & R. Soulier, (1984). Study of the seed oil from *A. spinosa* (L.), *Sapotaceae*. I. The glyceride fraction. *Revue française des Corps Gras*, **31**, 283-286.

Feucht W., P. P. S. Schmid & E. Christ. (1986). Distribution of flavonols in meristematic and mature tissues of *Prunus avium* shoots. *Journal of Plant Physiology*, **125**, 1-8.

Gershenzon J. (1984). Changes in the levels of plant secondary metabolites under water and nutrient stress. In: Phytochemical Adaptations to Stress (Timmermann B. N., Steelink C. and Loewus F. A., eds.), Plenum Press, New York, 273-320.

Huyghebaert A. & H. Hendrickx (1974). Quelques aspects chimiques physiques et technologiques de l'huile d'argan. *Oléagineux*, **1**, 29-31.

Keller M. & G. Hrazdina, (1998). Interaction of nitrogen availability during bloom and light intensity during veraison. II. Effects on anthocyanin and phenolic development during grape ripening. *American Journal of Enology and Viticulture*, **49**, 341-349.

Lois R., (1994). Accumulation of UV-absorbing flavonoids induced by UV-B radiation in *Arabidopsis thaliana* L. Part I. Mechanisms of UV-resistance in *Arabidopsis. Planta*, **194**, 498-503.

Maurin R., (1992). L'huile d'Argan : *Argania spinosa* (L.) Skeels *Sapotaceae*. *Revue française des Corps Gras*, **39**, 139-146.

Maurin R., K. Fellat-Zarrouk & R. Ksir, (1992). Positional analysis and determination of triacylglycerol structure of *Argania spinosa* seed oil. *Journal of the American Oil Chemist's Society*, **69**, 141-145.

Mc Murrough I. & J. Mc Dowell, (1978). Chromatographic separation and automated analysis of flavanols. *Analytical Biochemistry*, **91**, 92-100.

Neu R., (1956). Ein neues Reagenz zum Nachweis und zur Unterscheidung von Flavonen im Papierchromatogramm. Die *Naturwissenschaften*, **43**, 82.

Olsson L. C., M. Veit, G. Weissenböck & J. F. Borman, (1998). Differential flavonoid response to enhanced UV-B radiation in *Brassica napus*. *Phytochemistry*, **49**, 1021-1028.

Oulad-Ali A., V. Kirchner, A. Lobstein, B. Weniger & R. Anton, (1996). Structure elucidation of three triterpene glycosides from the trunk of *Argania spinosa*. *Journal of Natural Products*, **59**, 193-195.

Tahrouch S, Andary C., Rapior S., Mondolot L., Gargadennec A. and Fruchier A. (2000). Polyphenol investigation of *Argania spinosa* (Sapotaceae) endemic tree from Morocco. *Acta Botanica Gallica*, 147, (3), 225-232.

Tahrouch S., S. Rapior, J. M. Bessière & C. Andary, (1998). Volatile constituents of *Argania spinosa (Sapotaceae). Acta botanica Gallica*, **145**, 259-263.